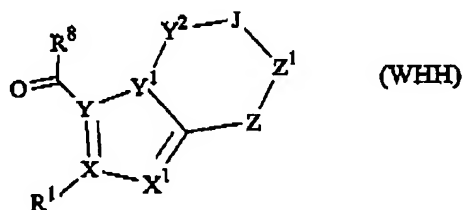


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OCT 04 2006AMENDMENTS TO THE CLAIMSThis listing of claims will replace all prior versions and listings of claims in the application.

1. (Previously presented) A compound of Formula (WHH)



wherein

R¹ is H, C₁₋₆alkyl, C₁₋₆haloalkyl, C₁₋₆alkoxy, C₁₋₆thioalkyl, cyano, halo, C₃₋₇cycloalkyl, -C₁₋₆alkylene-C₃₋₇cycloalkyl, C₂₋₆alkenyl or C₃₋₆alkynyl;

R⁸ is O-C₁₋₄alkyl, -N(CH₃)(OCH₃);

X is C;

Y is C;

X¹ is N;

Y¹ is N;

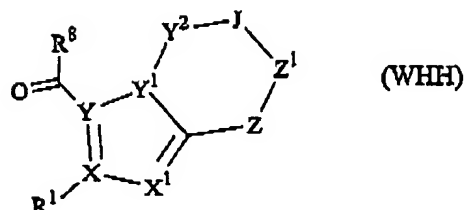
Y² is CH₂;

J is CH₂ or a bond;

Z¹ is CH₂ or C(O); and

Z is N-V, wherein V is phenyl, 2-pyridyl or 3-pyridyl substituted with two to three of the same or different substituents selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₆thioalkyl, C₁₋₄haloalkyl, halogen, N(C₁-C₄alkyl)₂ and CN.

2. (Previously presented) A process for preparing a compound of Formula (WHH)



wherein

R¹ is H, C₁₋₆alkyl, C₁₋₆haloalkyl, C₁₋₆alkoxy, C₁₋₆thioalkyl, cyano, halo, C₃₋₇cycloalkyl, -C₁₋₆alkylene-C₃₋₇cycloalkyl, C₂₋₆alkenyl or C₃₋₆alkynyl;
R⁸ is O-C₁₋₄alkyl, -N(CH₃)(OCH₃);

X is C;

Y is C;

X¹ is N;

Y¹ is N;

Y² is CH₂;

J is CH₂ or a bond;

Z¹ is CH₂ or C(O); and

Z is N-V, wherein V is phenyl, 2-pyridyl or 3-pyridyl substituted with two to three of the same or different substituents selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₆thioalkyl, C₁₋₄haloalkyl, halogen, N(C₁₋₄alkyl)₂ and CN;

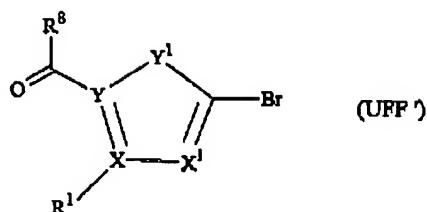
comprising reacting a compound of Formula (UFF)



wherein

Z, Z¹, J and Y² are defined as for Formula (WHH);

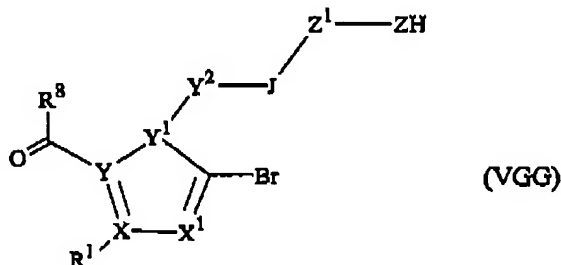
with a compound of Formula (UFF')

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wherein

 R^1 , R^8 , X, Y, X' and Y^1 are defined as for Formula (WHH);

in the presence of a suitable base and polar aprotic solvent to yield a compound of Formula (VGG)

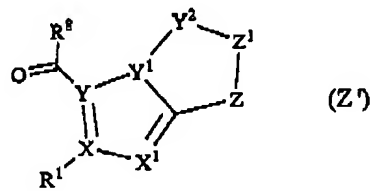
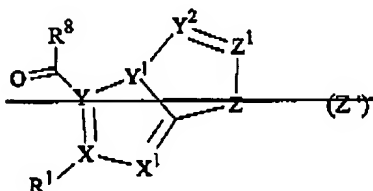


wherein

 R^1 , R^8 , X, Y, X' , Y^1 , Y^2 , J, Z^1 and Z are defined as for Formula (WHH);

and reacting said compound of Formula (VGG) with a high-boiling point polar aprotic solvent and a suitable silver salt under suitably high temperature.

3. (Currently Amended) A compound of Formula (Z')



wherein

R^1 is H, C_{1-6} alkyl, C_{1-6} haloalkyl, C_{1-6} alkoxy, C_{1-6} thioalkyl, cyano, halo, C_{3-7} cycloalkyl, $-C_{1-6}$ alkylene- C_{3-7} cycloalkyl, C_{2-6} alkenyl or C_{3-6} alkynyl;
 R^8 is $O-C_{1-4}$ alkyl, $-N(CH_3)(OCH_3)$;

X is C;

Y is C;

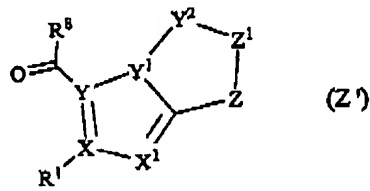
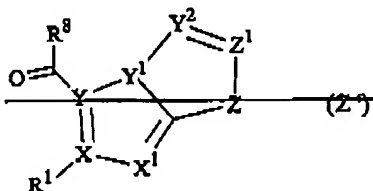
X¹ is N;Y¹ is N;Y² is CH or CR⁵;

R⁵ is selected from the group consisting of -CN, -C₁₋₄alk(en)ylene-CN, halo, C₁₋₆alkyl, C₂₋₆alkenyl, C₃₋₆alkynyl, C₁₋₆haloalkyl, aryl, -C₁₋₄alk(en)ylene-aryl, -C₁₋₄alk(en)ylene-heterocyclo, heterocyclo, -C₁₋₄alk(en)ylene-amino, -C₁₋₄alkylene-amino-C₁₋₄alkyl, aryl-amino, -amino-(C₁₋₆alk(en)yl)₁₋₂, -amino-aryl, -amino-heterocyclo, C₁₋₆alkoxy, -O-aryl and -O-heterocyclo;

Z¹ is C(O); and

Z is N-V, wherein V is phenyl, 2-pyridyl or 3-pyridyl substituted with two to three of the same or different substituents selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₆thioalkyl, C₁₋₄haloalkyl, halogen, N(C₁₋₄alkyl)₂ and CN.

4. (Currently Amended) A process for preparing a compound of Formula (Z')



wherein

R¹ is H, C₁₋₆alkyl, C₁₋₆haloalkyl, C₁₋₆alkoxy, C₁₋₆thioalkyl, cyano, halo, C₃₋₇cycloalkyl, -C₁₋₆alkylene-C₃₋₇cycloalkyl, C₂₋₆alkenyl or C₃₋₆alkynyl;
R⁸ is O-C₁₋₄alkyl, -N(CH₃)(OCH₃);

X is C;

Y is C;

X^1 is N;

Y^1 is N;

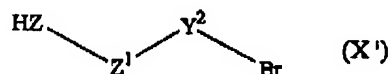
Y^2 is CH or CR⁵;

R^5 is selected from the group consisting of -CN, -C₁₋₄alk(en)ylene-CN, halo, C₁₋₆alkyl, C₂₋₆alkenyl, C₃₋₆alkynyl, C₁₋₆haloalkyl, aryl, -C₁₋₄alk(en)ylene-aryl, -C₁₋₄alk(en)ylene-heterocyclo, heterocyclo, -C₁₋₄alk(en)ylene-amino, -C₁₋₄alkylene-amino-C₁₋₄alkyl, aryl-amino, -amino-(C₁₋₆alk(en)yl)₁₋₂, -amino-aryl, -amino-heterocyclo, C₁₋₆alkoxy, -O-aryl and -O-heterocyclo;

Z^1 is C(O); and

Z is N-V, wherein V is phenyl, 2-pyridyl or 3-pyridyl substituted with two to three of the same or different substituents selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₆thioalkyl, C₁₋₄haloalkyl, halogen, N(C₁-C₄alkyl)₂ and CN;

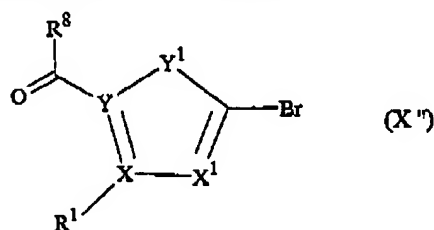
comprising reacting a compound of Formula (X')



wherein

Z , Z^1 and Y^2 are defined as for Formula (Z');

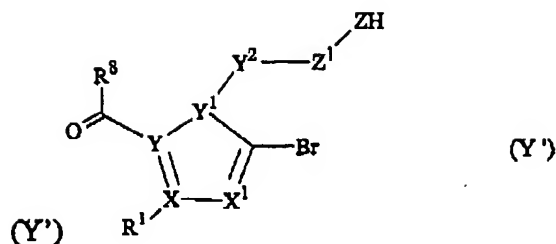
with a compound of Formula (UFF')



wherein

R^1 , R^8 , X , Y , X^1 and Y^1 are defined as for Formula (Z');

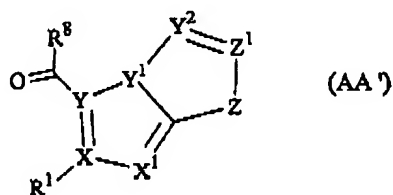
in the presence of a suitable base and polar aprotic solvent to yield a compound of Formula



wherein

R^1 , R^8 , X , Y , X^1 , Y^1 , Y^2 , Z^1 and Z are defined as for Formula (Z');
and reacting said compound of Formula (Y') with a high-boiling point polar aprotic solvent and a suitable silver salt under suitably high temperature.

5. (Previously Presented) A compound of Formula (AA')



wherein

R^1 is H, C_{1-6} alkyl, C_{1-6} haloalkyl, C_{1-6} alkoxy, C_{1-6} thioalkyl, cyano, halo, C_{3-7} cycloalkyl, $-C_{1-6}$ alkylene- C_{3-7} cycloalkyl, C_{2-6} alkenyl or C_{3-6} alkynyl;

R^8 is $O-C_{1-4}$ alkyl, $-N(CH_3)(OCH_3)$;

X is C;

Y is C;

X^1 is N;

Y^1 is N;

Y^2 is CH or CR^5 ;

R^5 is selected from the group consisting of $-CN$, $-C_{1-4}alk(en)ylene-$
 CN , halo, $C_{1-6}alkyl$, $C_{2-6}alkenyl$, $C_{3-6}alkynyl$, $C_{1-6}haloalkyl$, aryl,
 $-C_{1-4}alk(en)ylene-aryl$, $-C_{1-4}alk(en)ylene-heterocyclo$, heterocyclo,
 $-C_{1-4}alk(en)ylene-amino$, $-C_{1-4}alkylene-amino-C_{1-4}alkyl$, aryl-

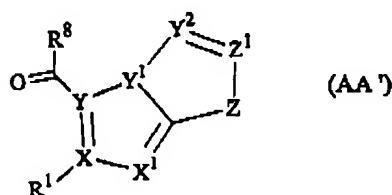
amino, -amino-(C₁₋₆ alk(en)yl)₁₋₂, -amino-aryl, -amino-heterocyclo,
C₁₋₆alkoxy, -O-aryl and -O-heterocyclo;

Z¹ is CR⁷;

wherein R⁷ is chloro or bromo; and

Z is N-V, wherein V is phenyl, 2-pyridyl or 3-pyridyl substituted with two to three of the same or different substituents selected from the group consisting of C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₆thioalkyl, C₁₋₄ haloalkyl, halogen, N(C₁-C₄alkyl)₂ and CN.

6. (Currently Amended) A process for preparing a compound of Formula (AA')



wherein

R¹ is H, C₁₋₆alkyl, C₁₋₆haloalkyl, C₁₋₆alkoxy, C₁₋₆thioalkyl, cyano, halo, C₃₋₇cycloalkyl, -C₁₋₆alkylene-C₃₋₇cycloalkyl, C₂₋₆alkenyl or C₃₋₆alkynyl;

R⁸ is O-C₁₋₄alkyl, -N(CH₃)(OCH₃);

X is C;

Y is C;

X¹ is N;

Y¹ is N;

Y² is CH or CR⁵;

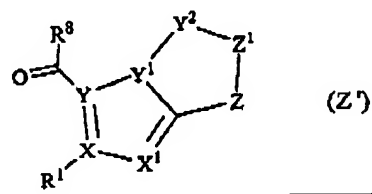
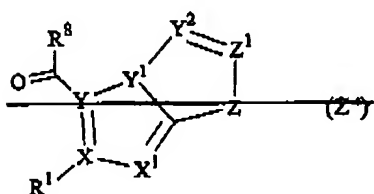
R⁵ is selected from the group consisting of -CN, -C₁₋₄alk(en)ylene-CN, halo, C₁₋₆alkyl, C₂₋₆alkenyl, C₃₋₆alkynyl, C₁₋₆haloalkyl, aryl, -C₁₋₄alk(en)ylene-aryl, -C₁₋₄alk(en)ylene-heterocyclo, heterocyclo, C₁₋₄alk(en)ylene-amino, -C₁₋₄alkylene-amino-C₁₋₄alkyl, aryl-amino, -amino-(C₁₋₆ alk(en)yl)₁₋₂, -amino-aryl, -amino-heterocyclo, C₁₋₆alkoxy, -O-aryl and -O-heterocyclo;

Z^1 is CR^7 ;

wherein R^7 is chloro or bromo; and

Z is N-V, wherein V is phenyl, 2-pyridyl or 3-pyridyl substituted with two to three of the same or different substituents selected from the group consisting of C_{1-4} alkyl, C_{1-4} alkoxy, C_{1-6} thioalkyl, C_{1-4} haloalkyl, halogen, $N(C_{1-4}alkyl)_2$ and CN;

comprising reacting a compound of Formula (Z')



wherein

R^1 , R^8 , X, Y, X^1 , Y^1 , Y^2 , and Z are defined as for Formula (AA'); and

Z^1 is C(O);

with phosphoryl trichloride or phosphoryl tribromide, neat or with a suitable solvent and without a base or with a suitable base.